

## **REMARKS**

I. Regarding claim 24, the first full paragraph on page 2 of the instant application teaches that an antagonist of interest reduces ghrelin activity (is a ghrelin antagonist) and reduces GHSR 1a activity (is a GHSR 1a antagonist); the paragraph bridging pages 1 and 2 teaches the functions associated with a ghrelin or a GHSR 1a; and in the paragraph bridging pages 13 and 14 teaches the need and/or benefit of reducing ghrelin or GHSR 1a function in certain conditions. Regarding claim 29, the first full paragraph of page 15 teaches competition assays to find ghrelin antagonists, using assays known in the art, wherein, for example, ghrelin is labeled. Accordingly, no issue of new matter arises with the above amendments, the amendments place the claims in condition for allowance and at the least, the amendments simplify issues for appeal. Hence, entry of the amendments is requested respectfully.

II. At the lower half of page 2 of the Office Action, the Examiner raised an objection to claims 2, 6, 12, 13 and 31-36 alleging that the claims were essentially duplicates of an allowed claim. Claims 14 and 37-41 were alleged not to limit further a prior claim.

The objection is traversed for the following reasons.

A nucleic acid which has the sequence of SEQ ID NO:8 as recited in claim 1 is one which contains at least SEQ ID NO:8, see, for example, the last full paragraph on page 8, the third full paragraph on page 11, the paragraph bridging pages 14 and 15, Examples 3 and 4, and Figures 22-25, which teach that the nucleic acid of interest can contain, for example, flanking sequences and retain the desired activity.

Hence, a nucleic acid of interest can bind to different molecules, can have varying overall structure and can have varying properties without deviating from the function of SEQ ID NO:8, that is, antagonizing ghrelin or GHSR 1a function, for example, by having varying length or varying affinity levels for a ligand.

In any event, to advance prosecution and to place the instant application in condition for allowance, applicants canceled claims 2, 12-14 and 31-41 without prejudice to prosecution in a continuation application. Claim 6 was amended to further define the ghrelin structure. Accordingly, the objection can be withdrawn.

III. At the bottom of page 4 of the Office Action, claims 16-18, 20-22 and 29 were rejected under 35 U.S.C. 112, second paragraph.

The rejection is traversed for the following reasons.

Claims 16 and 20 recite methods to obtain the nucleic acid of claim 1. Claim 29 relates to a competition assay, essentially as known in the art. Hence, the antagonist of claim 1 can be used to locate other molecules which also antagonize ghrelin or GHSR 1a function in a competition assay.

In view of the claim amendments, withdrawal of the rejection is in order.

IV. At the top of page 6 of the Office Action, claims 24 and 25 were rejected under 35 U.S.C. 112, first paragraph for an alleged want of enablement.

The rejection is traversed for the following reasons.

The instant specification teaches ghrelin regulating, for example, appetite, see paragraph bridging pages 1 and 2. Examples 8, 9 and 12 show the expected *in vivo* effect of an antagonist of interest. Example 11 shows a reduction in growth hormone release on expression of an antagonist of interest.

The molecule of interest can be configured to be stable to naturally occurring nucleases, for example, by containing an L-base. Hence, a molecule of interest will have enhanced plasma half life.

Submitted herewith is a copy of Shearman et al., *Endo* 147(3):1517-1526, 2006, which reports on studies conducted using antagonists of interest. As provided in the instant specification, the antagonists of Shearman et al. blocked ghrelin-induced feeding, promoted weight loss and reduced fat mass.

Accordingly, the antagonist of interest functions as taught in the instant specification *in vitro* and *in vivo*. The instant specification teaches how to make and how to use the invention of interest.

As noted on page 1523, right column, first full paragraph of Shearman et al., another study using ghrelin knockout mice observed the opposite phenotype as reported in Sun et al. of

record. Hence, the working examples of the instant specification and Shearman et al. support the claimed invention.

In view thereof, the instant specification enables the claimed invention. Accordingly, the rejection can be removed.

V. Claims 1, 27 and 45 were allowed. Claims 3, 4, 9-11, 26 and 30 are allowable if rewritten in independent form including limitations of the base claim and any intervening claims.

Claims 3, 4, 26 and 30 depend on allowed claim 1. Claims 9 and 10 were canceled without prejudice. Claim 6 was amended to exemplify an element.

## **CONCLUSION**

Applicants have taken steps to place the instant application in condition for allowance. Reexamination, reconsideration, withdrawal of the objection and rejections, and early indication of allowance are solicited earnestly.

Respectfully submitted,

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